



Sughrue

SUGHRUE MION ZINN MACPEAK & SEAS, PLLC

2000 RECEIVED PTO
09/913611

11 6 AUG 2001

2100 Pennsylvania Avenue, NW
Washington, DC 20037-3213

T 202.293.7060
F 202.293.7860

www.sughrue.com

August 16, 2001

BOX PCT

Commissioner for Patents
Washington, D.C. 20231

PCT/JP00/00875

-filed February 16, 2000

Re: Application of Shigeru MURAKAMI, Yoichi NAMBA, Tsutomu MASUKO and Yuichi KAMIJO
CARBON FIBER WOVEN FABRIC AND METHOD FOR PRODUCTION THEREOF
Assignee: **SHOWA DENKO K.K.**
Our Ref: Q54917

Dear Sir:

The following documents and fees are submitted herewith in connection with the above application for the purpose of entering the National stage under 35 U.S.C. § 371 and in accordance with Chapter II of the Patent Cooperation Treaty:

- ☒ an executed Declaration and Power of Attorney.
- ☒ an English translation of the International Application.
- ☒ one (1) sheet of drawings.
- ☒ an English translation of Article 34 amendments.
- ☒ International Preliminary Examination Report (foreign language)
- ☒ an executed Assignment and PTO 1595 form.
- ☒ International Search Report and a Form PTO-1449.
- ☒ a Preliminary Amendment

It is assumed that copies of the International Application, and any Articles 19 and 34 amendments as required by § 371(c) will be supplied directly by the International Bureau, but if further copies are needed, the undersigned can easily provide them upon request.

The Government filing fee is calculated as follows:

Total claims	12	-	20	=		x	\$18.00	=	\$0.00
Independent claims	1	-	3	=		x	\$80.00	=	\$0.00
Base Fee									\$860.00

TOTAL FILING FEE

\$860.00

Recordation of Assignment

\$ 40.00

TOTAL FEE

\$900.00

Checks for the statutory filing fee of \$860.00 and Assignment recordation fee of \$40.00 are attached. You are also directed and authorized to charge or credit any difference or overpayment to Deposit Account No. 19-4880. The Commissioner is hereby authorized to charge any fees under 37 C.F.R. §§ 1.16, 1.17 and 1.492 which may be required during the entire pendency of the application to Deposit Account No. 19-4880. A duplicate copy of this transmittal letter is attached.

Priority is claimed from February 18, 1999, and June 9, 1999, based on Japanese Application Nos. 11-40518 (Pat. Appln.), and 11-161756 (Pat. Appln.). Benefit is claimed from April 7, 1999 and June 30, 1999 based on U.S. Provisional Application Nos. 60/128,054 and 60/141,244.

Respectfully submitted,

Bruce E. Kramer

Bruce E. Kramer

Registration No. 33,725

BEK/amt

09/913611
531 Rec'd PCT. 16 AUG 2001

PATENT APPLICATION

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re application of

Shigeru MURAKAMI, et al.

Appln. No.: Not Yet Assigned

Group Art Unit: Not Yet Assigned

Confirmation No.: Not Yet Assigned

Examiner: Not Yet Assigned

Filed: August 16, 2001

For: CARBON FIBER WOVEN FABRIC AND METHOD FOR PRODUCTION THEREOF

PRELIMINARY AMENDMENT

Commissioner for Patents
Washington, D.C. 20231

Sir:

Prior to examination, please amend the above-identified application as follows:

IN THE CLAIMS:

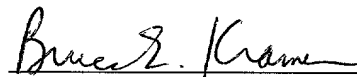
Please enter the following amended claim:

1. A carbon fiber woven fabric characterized by being obtained by firing a cellulose-based woven fabric, and by having a thickness in the range of 0.05-0.4 mm, a volume resistivity of less than 0.2 $\Omega \cdot \text{cm}$ in the layer direction, and a gas permeability of not less than 1500 cc/cm²/hr/mmAq.

REMARKS

Claim 1 has been amended based on the disclosure at page 4, line 36 in the present application. Entry and consideration of this Amendment is respectfully requested.

Respectfully submitted,



Bruce E. Kramer
Registration No. 33,725

SUGHRUE, MION, ZINN,
MACPEAK & SEAS, PLLC
2100 Pennsylvania Avenue, N.W.
Washington, D.C. 20037-3213
Telephone: (202) 293-7060
Facsimile: (202) 293-7860
BEK/amt
Date: August 16, 2001

09/913611

531 Rec'd PCT: 16 AUG 2001

AMENDMENT

Attorney Docket No. Q54917

APPENDIX

VERSION WITH MARKINGS TO SHOW CHANGES MADE

IN THE CLAIMS:

The claims are amended as follows:

1. A carbon fiber woven fabric characterized by being obtained by firing a cellulose-based woven fabric, and by having a thickness in the range of 0.05-0.4 mm, a volume resistivity of ~~not~~ less than 0.2 $\Omega \cdot \text{cm}$ in the layer direction, and a gas permeability of not less than 1500 cc/cm²/hr/mmAq.



#5
JC07 Rec'd PCT/PTO 19 MAR 2002

09/913611

Attorney Docket No. Q54917
PATENT APPLICATION

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re application of

Shigeru MURAKAMI, et al.

Appln. No. 09/913,611

Group Art Unit: Not Yet Assigned

Confirmation No.: Not Yet Assigned

Examiner: Not Yet Assigned

Filed: August 16, 2001

For: CARBON FIBER WOVEN FABRIC AND METHOD FOR PRODUCTION THEREOF

RESPONSE TO NOTIFICATION OF DEFECTIVE RESPONSE

Commissioner for Patents
Washington, D.C. 20231

Sir:

In response to the Notification of Defective Response, dated February 19, 2002, a copy of which is enclosed, and the Notification of Missing Requirements Under 35 USC § 371, dated November 27, 2001, a copy of which is enclosed, applicants submit that the Declaration filed by applications on August 16, 2001, a copy of which is attached, complies with 37 C.F.R. § 1.497(a) and (b).

The Notification of Defective Response lists two alleged errors in the Declaration.

The first alleged error is that the name of the third inventor is spelled differently on the Declaration than on the 1A. Applicants point out, however, that if the Japanese characters of a name are shown in Roman letters, the third inventors' name is usually indicated in one of two ways, that is, either "Tsutomu Masuko" or "Tutomu Masuko". The characters of "Tsu" can read the same as the characters of "Tu" in Japanese. This means that these two names are considered

RESPONSE TO NOTIFICATION OF DEFECTIVE RESPONSE
U.S. Appln. No. 09/913,611

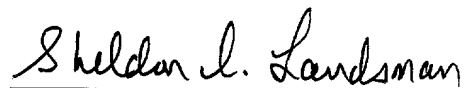
to be the same in Japan. Mr. Masuko signed the Declaration with the spelling he prefers "Tsutomo". Applicants submit that the spelling of the third named inventor on the Declaration, therefore, is correct and proper.

The second alleged error is that the Declaration lists a fourth inventor, but the fourth inventor is not listed on the 1A. Applicants point out, however, that the fourth inventor, Mr. Yuichi Kamijo, was added at the international stage. Applicants have received a Notification of the Recording of a Change, dated August 27, 2001, from the International Bureau, a copy of which is enclosed, showing that the name of the fourth inventor, Mr. Yuichi Kamijo, has been added at the international stage. Thus, the name of the fourth inventor does not appear on the 1A, but the inclusion of the fourth inventor on the Declaration is proper and necessary.

Therefore, no correction of the Declaration is needed.

In view of the above, applicants submit that the Notification of Defective Response and Notification of Missing Requirements Under 35 U.S.C. § 371 were issued in error, and that no Declaration or fee need to be filed in response to the Notification and Notice. If it is determined that a fee is needed, or a petition for extension of time or any other petition, please charge the fee to our Deposit Account No. 19-14480 and treat this paper as a petition.

Respectfully submitted,



Sheldon I. Landsman
Registration No. 25,430

SUGHRUE MION, PLLC
2100 Pennsylvania Avenue, N.W.
Washington, D.C. 20037-3213
Telephone: (202) 293-7060
Facsimile: (202) 293-7860

Date: March 19, 2002

1/PRTS

09/913611

SHD-H808

- 1 -

531 Rec'd PCT/

16 AUG 2001

SPECIFICATION

CARBON FIBER WOVEN FABRIC AND METHOD FOR PRODUCTION THEREOF

5

Cross Reference to Related Applications

This application claims the benefit of priority
right based on U.S. Provisional Application No.
60/128054, filed on April 7, 1999 and U.S. Provisional
10 Application No. 60/141244, filed on June 30, 1999.

Technical Field

The present invention relates to a carbon fiber
woven fabric manufactured by firing a cellulose-based
15 woven fabric, and to a process for its manufacture. The
carbon fiber woven fabric has excellent chemical
resistance, electric conductivity, heat conductivity, gas
permeability and handleability and is a thin porous
carbon sheet with uniform gas permeability within its
20 layer, and it is therefore useful as a gas diffusing
porous carbon sheet, catalyst sheet or the like,
particularly for solid polymer fuel cells.

Background Art

25 Electric vehicles (EVs) and fuel cells have been
developed in recent years in response to the problem of
environmental pollution by exhaust gas from internal
combustion engines of automobiles and the like, and
further development of fuel cells has become essential in
30 light of the problems of range, total energy efficiency
and infrastructure which includes electrical recharging
equipment; it has therefore been a goal to achieve fuel
cells with higher performance and lighter weight.

Various types of fuel cells have been developed,
35 such as alkali types, phosphoric acid types, melted
carbonate types, solid electrolyte types and solid
polymer types, depending on the nature of the electrolyte

solution used, but attention is being focused on solid polymer type fuel cells as power sources because of their operability at low temperature, easy handleability and high output density.

5 The important components of a solid polymer fuel cell are the ion-exchange membrane to maintain appropriate moisture, and the conductive porous diffusion sheet for permeation treatment of the water and gas generated during the cell reaction. The diffusion sheet
10 is usually a porous carbon thin sheet. (Hereunder this will be referred to as a "porous carbon sheet".)

 The properties required for gas diffusion porous carbon sheets that are used in solid polymer fuel cells include thinness, satisfactory gas permeability, uniform
15 gas permeability within the sheet layer, open pores in the porous sheet, excellent heat and electrical conductivity, low volume resistivity in the thickness direction, suitable strength for incorporation into the electrolyte cell, and excellent handleability.

20 Fuel cell gas diffusion porous carbon sheets hitherto disclosed include the porous carbon sheets for fuel cells described in Japanese Unexamined Patent Publication HEI No. 3-285873 and Japanese Unexamined Patent Publication HEI No. 5-254957, wherein carbon
25 fibers are impregnated with a thermosetting resin as a binder, and firing and carbonization are carried out to manufacture a porous electrolyte sheet for fuel cells having numerous fine pores in the direction perpendicular to the sheet surface, and a carbon sheet with improved
30 air permeability, heat conductivity, compressive strength, etc.

 In addition, as in Japanese Examined Patent Publication HEI No. 2-58369, HEI No. 2-23505 and HEI No. 7-115970, for example, it has been disclosed that the
35 most inexpensive manufacturing processes for such carbon sheets involve paper making of organic fibers and cellulose, and some of these methods have been tested.

However, because these methods require impregnation of thermosetting resins and high temperature carbonizing firing treatment, they have been characterized by poor productivity and high cost.

5 With recent advances in development, there is, in particular, a demand for even lower cost and high performance for solid polymer fuel cells, while lower cost and high performance are also becoming essential for gas diffusion porous carbon sheets. Moreover, because of
10 the smaller size and lighter weights, thinner sheets on the order of no more than a few hundred micrometers have been desired.

15 The following fundamental problems are associated with porous carbon sheets obtained by the paper making methods mentioned above.

20 1) Because the organic fibers are made into paper sheets, the fibers are horizontal during the paper making process. That is, since the fiber orientation is aligned in the edgewise or layer direction, the heat and electrical conductivity in the layer transverse direction
25 are inferior. It is therefore necessary to add conductive carbon particles during paper making process to increase the electrical conductivity, and this results in inconveniences such as reduced mechanical strength and gas permeability.

30 2) Because the prepared paper-like sheet is passed through a step of soaking in a resin solution, numerous closed holes with no effective function are present in the porous carbon sheet. The gas permeability is therefore not improved.

35 3) When creating a thin product, the starting paper-made sheet must also be thin, and this makes manufacturing difficult by the paper making method. This is because it is very difficult to maintain uniformity within the layer of the paper-made sheet, and the gas permeability within the layer also becomes non-uniform.

4) Porous carbon sheets prepared by soaking paper-

made sheets in resins and heat treating them have low elasticity and are hard and fragile and, therefore, exhibit poor handleability during the assembly of cells.

5 Because the paper-making methods of the prior art are problematic in meeting these demands, new improved manufacturing methods have been desired.

10 Moreover, fuel cells generate large amounts of water during operation, due to their nature. When the water spills into the cell reaction zone, it creates a state of flooding, where the oxygen or other reaction gas supply channel becomes cut off and the cell reaction no longer takes place, resulting in drastically reduced cell output. Therefore porous carbon sheets with some degree of water repellency have also been desired.

15 Methods of weaving carbon fibers into woven fabrics have been considered in the past, but such methods are hampered by the high cost of carbon fibers, the steps for weaving, and the fact that the woven fabric holds the fibers in a horizontal state such that the properties in
20 the layer transverse direction are unsatisfactory, as mentioned above.

Disclosure of the Invention

25 As a result of diligent study, the present inventors have considered the use of cellulose-based fiber woven fabrics as starting materials, and have attempted to use them for fired and carbonized carbon fiber woven fabrics. It was found that inexpensive carbon fiber woven fabrics with relatively good gas permeability can be obtained
30 thereby.

The present invention therefore provides the following:

35 (1) A carbon fiber woven fabric characterized by being obtained by firing a cellulose-based woven fabric, and by having a thickness in the range of 0.05-0.4 mm, a volume resistivity of less than 0.2 $\Omega \cdot \text{cm}$ in the layer direction, and a gas permeability of not less than 1500

cc/cm²/hr/mmAq.

(2) A carbon fiber woven fabric according to (1), wherein the compressive strength is not less than 70 kgf/cm².

5 (3) A carbon fiber woven fabric according to (1) or (2), wherein the electrical resistance in the direction of thickness of the woven fabric is no greater than 50 mΩ·cm² as measured between two copper plates with a load of 4 kgf/cm².

10 (4) A carbon fiber woven fabric according to (1) to (3), wherein the orientation of the carbon fiber woven fabric as defined in the present specification includes an orientation component of 4/9 or greater.

15 (5) A carbon fiber woven fabric according to (1) to (4), wherein the orientation of the carbon fiber woven fabric as defined in the present specification is an average of 1/3 or greater.

(6) A carbon fiber woven fabric according to (1) to (5) which is a plain weave.

20 (7) A carbon fiber woven fabric according to (1) to (6) which has a water repellent property.

(8) A gas diffusion porous carbon sheet for a solid polymer fuel cell which comprises a carbon fiber woven fabric according to (1) to (7).

25 (9) A process for the manufacture of a carbon fiber woven fabric, characterized by firing a cellulose-based woven fabric in a non-oxidizing atmosphere.

(10) A process for the manufacture of a carbon fiber woven fabric according to (9), wherein the cellulose-based woven fabric is soaked with a phosphoric acid or phosphorus compound solution.

30 (11) A process for the manufacture of a carbon fiber woven fabric according to (9) or (10), wherein the firing temperature is in the range of 900-3000°C.

35 (12) A process for the manufacture of a carbon fiber woven fabric according to (9) to (11), which includes coating the fired carbon fiber woven fabric with a water-

repellent resin.

Brief Description of the Drawings

5 Figs. 1A to 1C show plain woven fabrics, wherein Fig. 1A is a plan view, Figs. 1B and 1C are lengthwise and widthwise cross-sectional views in the directions of X and Y, and Fig. 1C illustrates the definition of the degree of orientation according to the invention.

10 Fig. 2 is a cross-sectional view of the basic construction of a solid polymer fuel cell using a porous carbon sheet of the invention.

Best Mode for Carrying Out the Invention

15 The present invention is characterized in that for manufacture of a porous carbon sheet, a cellulose-based fiber woven fabric is fired as the starting material.

According to the invention, the carbon fiber sheet is prepared as a fabric so that despite its thinness it can exhibit sufficient strength and excellent uniformity within its surface.

20 Furthermore, when a fabric of the starting fibers before carbonization-firing is used instead of using carbon fibers and weaving them into a woven fabric, the starting fiber yarn itself which has free orientation properties is soft with the fibers not aligned in the edgewise or layer direction, as opposed to a stiff woven fabric of carbon fibers, and therefore the orientation property is greater in the layer transverse direction and the electrical conductivity, heat conductivity and compressive strength in the thickness direction are superior.

30 Moreover, the carbon fiber woven fabric of the invention has a high open void volume with no closed voids, since there is no step of soaking in a resin solution as with the paper making methods described above. The cellulose-based woven fabric used as the starting material is subjected to solid phase

35

carbonization without melting, and therefore the form of the fibers of the green fabric before firing carbonization reflects the carbon fiber woven fabric as the finished product, so that there is no breakage of the fibers, the strength is excellent and the handleability is satisfactory.

Japanese Examined Patent Publication HEI No. 2-58369 describes improvement in strength by impregnation treatment in a metal phosphate, but sheets prepared by paper making methods such as the method disclosed here have short and non-continuous carbon fibers or else the fibers are horizontal, for which reasons the improved strength of the porous carbon sheet is not reflected to any great extent; however, it is believed that when the filaments are continuous or not many of the components are oriented in the direction of thickness, such as in the carbon fiber woven fabric of the invention, there is exhibited a greater effect of improvement in the strength by the phosphoric acid or phosphorus compound.

After filling the carbon fiber woven fabric of the invention with resin powder, it may be heat treated at the resin melting temperature to achieve improved strength by the resin reinforcement and to exhibit water repellency. Since the fibers are horizontal in the case of paper making or in the case of a woven fabric of carbon fibers, it is difficult to uniformly fill the interior of the sheet with the resin powder. There is a slight effect of impregnation of the resin after firing-carbonization on the gas permeability, but it is not enough to impair the properties to any great extent, unlike impregnation of the resin before firing, as in paper making methods due to differences in the resin wettability leading to different permeability of the resin into the fabric.

The fabric composed of cellulose-based fibers or cellulose-containing fibers as the starting material according to the invention may be a natural cellulose

fiber fabric of cotton, hemp or the like, or it may be an artificial cellulose fiber fabric of viscose artificial silk, acetate artificial silk, or the like.

5 The method of weaving the starting fabric is not particularly restricted and the weave may be any of various types including a plain weave, twill weave or satin weave, but it is preferably a plain weave from the standpoints of uniformity within the surface and of strength.

10 For example, a plain woven fabric is very inexpensive and readily obtainable, as well as quite easy to handle. The starting material for the invention is an already woven fabric, and a fabric with a large area may be cut out beforehand depending on the intended use for
15 provision as the starting material in the manufacturing process of the invention, or a carbon fiber woven fabric manufactured with a large area may be cut out to the necessary size.

20 The steps are greatly simplified since there is no step of weaving carbon fibers, as for conventional carbon fiber woven fabrics.

According to the invention, it was found that a cotton fabric obtained particularly by plain weaving of cellulose-based fiber spun yarn or twisted yarn, or yarn
25 obtained using fibers of no greater than a specified thickness, can be used as a starting material to give a carbon fiber woven fabric with excellent properties for fuel cells.

30 The carbon fiber woven fabric of the invention is characterized in that by using a cellulose-based fabric as the starting material before firing, since the starting fiber yarn itself has a soft nature, the orientation in the layer transverse direction of the cellulose-based fiber woven fabric, and therefore of the
35 carbon fiber woven fabric, (the direction normal to the surface or layer direction of the woven fabric) can be increased.

According to the invention, the orientation in the direction of the thickness of the carbon fiber woven fabric is defined as follows.

5 Figs. 1A to 1C are cross-sectional illustrations of carbon fiber fabrics obtained with plain woven fabrics as the starting materials. The carbon fiber woven fabric of the invention retains virtually the exact shape of the cellulose-based fibers. Thus even a cross-sectional view shows virtually the same cross-sectional shape as the
10 cellulose-based fibers. The fibers are sewn with the weft aligned at equal spacings and the warp woven therein; here, p is the horizontal component and q the vertical component of the line L connecting the lowest point A and highest point B of the center of the warp
15 from one weft to the next weft. The value of p is equal to the weft spacing. If the fibers are free of narrowing or crushing distortion and the weaving is carried out as shown in Figs. 1A to 1C, then q may be considered almost the value of the fabric width w minus the warp thickness
20 d .

The orientation is defined as $q/(p + q)$. According to this definition, the orientation is 0 when completely horizontal and 1 when completely vertical, although in actuality intermediate values are exhibited with
25 continuous fibers.

The orientation component can be measured by observing the cross-section and yarn thickness with a light microscope, an SEM or the like.

30 In the case of fabrics such as plain weaves in which the entirety is uniformly woven, the average orientation of the total fabric can be determined by examining one section thereof.

The carbon fiber woven fabric of the invention, reflecting the cellulose-based woven fabric which is used
35 as the starting material, may have a high orientation of $1/3$ or greater or even $4/9$ or greater at certain sections of the woven fabric. By thus including components with

high orientation (referred to as orienting components) in all or a part of the carbon fiber woven fabric, it is possible to improve the electric conductivity and heat conductivity in the layer transverse direction of the entire woven fabric. Orienting components with a higher degree of orientation are therefore preferred.

When the woven fabric is a plain weave, the orientation of each orienting component is almost exactly the average orientation of the entire carbon fiber woven fabric, but with special fabric textures other than plain weaves, such as diagonal weaves or satin weaves, the orientation of specific orienting components are not necessarily the average orientation of the woven fabric as a whole. The carbon fiber woven fabric of the invention can exhibit a high value even for the average orientation, compared to mechanically woven carbon fibers or pressed split carbon fibers obtained using conventional carbon fibers. The upper limit for the average orientation is theoretically $1/2$ when using filaments having the same thickness for warp and weft in the absence of narrowing or crushing of the fibers, but if the warp and weft thickness are altered a value of greater than $1/2$ is also possible. By thus having a high average orientation, the carbon fiber woven fabric of the invention can exhibit high electric conductivity and high heat conductivity in the layer transverse direction.

According to the invention, the cellulose-based woven fabric yarn used as the starting material is preferably constructed with fibers having a post-firing thickness of no greater than $10\text{ }\mu\text{m}$. By using filament-forming fibers of such small thickness, it is possible to achieve more open pores in-plane contributing to the gas permeability and thus obtain a higher performance gas diffusing sheet. However, if the fibers are too thin the gas permeability is affected by the exponential factor of the open pores size, leading to inconveniences such as lower gas permeability.

According to the invention, this manner of cellulose-based woven fabric as the starting material may be fired in a non-oxidizing atmosphere to produce a carbon fiber woven fabric having properties including a
5 thickness of 0.05-0.4 mm, a compressive strength of 70 kgf/cm² or greater and a gas permeability of 1500 cc/cm²/hr/mmAq or greater.

The thickness of the carbon fiber woven fabric of the invention is in the range of 0.05-0.4 mm. If the
10 thickness is less than 0.05 mm, the strength is insufficient for the demands of a gas diffusing carbon sheet used in a solid polymer-type fuel cell, and if it is greater than 0.4 mm, the thickness of cell stacks in solid polymer-type fuel cells increases to an impractical
15 level.

The thickness of the woven fabric of the starting material for obtaining a carbon fiber woven fabric of the desired thickness will usually need to be between 0.1 and 0.2 mm thicker. Selection of the starting material
20 thickness must be confirmed by prior testing under the same conditions. The thickness of the starting material woven fabric thickness (filament thickness) can be selected to give a carbon fiber woven fabric having a thickness of 0.05-0.4 mm according to the invention.

The compressive strength of the carbon fiber woven fabric of the invention is 70 kgf/cm² or greater. Here, the compressive strength is the value obtained by laminating 50 samples, compressing at a loading rate of 0.5 mm/min, measuring the maximum load resistance,
25 dividing this by the area, and calculating the maximum load resistance per unit area. If the compressive strength is less than 70 kgf/cm², the sheet will break when clamped during assembly of a cell stack.

The gas permeability of the carbon fiber woven fabric of the invention is 1500 cc/cm²/hr/mmAq or
35 greater. The gas permeability is the value obtained by calculation from the volume (cc) of permeating gas per

hour (hr) and the differential pressure P (water pressure, mmAq), based on the following equation, for air passing at 3000 mL/min through a 50 cm² carbon fiber fabric.

5 Gas permeability = $3000 \text{ mL/min} \times 60 \text{ min} / 50 \text{ cm}^2 / P \text{ mmAq}$
If the gas permeability is less than 1500 cc/cm²/hr/mmAq, the fuel cell electric generating properties are impaired. The gas permeability is preferably 3000 cc/cm²/hr/mmAq or greater.

10 The carbon fiber woven fabric of the invention may have a volume resistivity of less than 0.2 Ω·cm and even not more than 0.13 Ω·cm in the planar direction.

The carbon fiber woven fabric of the invention may have an electric resistance of no greater than 50 m·Ω·cm²
15 and even no greater than 40 m·Ω·cm² in the layer transverse direction, as measured between copper plates with a load of 4 kgf/cm². As will be clear from the aforementioned method of measurement, this resistance in the layer transverse direction includes the contact
20 resistance with the copper plates, but since the carbon fiber woven fabric of the invention is characterized by the above-mentioned high orientation in the layer transverse direction, it will have a low electrical resistance in the layer transverse direction.

25 The carbonization rate of the carbon fiber woven fabric of the invention can be greatly increased by soaking the cellulose-based woven fabric in phosphoric acid or a phosphoric acid compound before firing.

The carbon fiber woven fabric of the invention may
30 also be imparted with water repellency. Water repellency can be achieved by coating the carbon fiber woven fabric with a resin. Such coating with a resin will have also have an effect of enhancing the strength of the carbon fiber woven fabric. The resin used may be any resin that
35 exhibits water repellency and thermoplasticity, such as polyvinylidene fluoride, divinylbenzene, Teflon (TM),

polyethylene, polypropylene or the like. Fluorine-containing resins are preferred for their high water repellency.

5 The amount of resin may be in the range of 5-60 wt% based on the carbon fiber woven fabric. At less than 5 wt% the imparted water repellency is insufficient, while at greater than 60 wt% properties including electrical resistance and gas permeability may be undesirably affected.

10 A process for the manufacture of a carbon fiber woven fabric of the invention will now be explained.

15 According to the invention, the carbon fiber woven fabric may be prepared by soaking the cellulose-based woven fabric in a phosphoric acid or phosphoric acid compound solution if necessary, and then firing in a non-oxidizing atmosphere.

20 The phosphoric acid or phosphoric acid compound used in the process of the invention to provide an effect of improving the carbonization rate includes aqueous phosphoric acid as well as phosphates such as aqueous ammonium phosphate, magnesium phosphate, calcium phosphate, sodium phosphate, potassium phosphate and aluminum phosphate, among which water-soluble aqueous aluminum phosphate is preferred for ease of handleability and operation.

25 A coating method, an immersion method or the like may be employed to soak the cellulose-based fiber woven fabric with the phosphoric acid or phosphoric acid compound.

30 The firing and carbonization are carried out, for example, by firing in a non-oxidizing atmosphere in coke in an Atchison furnace or in a Reed Hammer firing furnace, or in a non-oxidizing atmosphere of an inert gas such as nitrogen or argon.

35 The firing temperature is preferably 900°C or higher in order to obtain strength characteristics which allow use as porous carbon sheets for fuel cells.

In special consideration of the electric conductivity, the firing is more preferably carried out at 1500°C to 2300°C. It is preferably no higher than 3000°C because with higher temperatures the strength
5 begins to lower and the gas permeation is reduced.

The fired and carbonized carbon fiber woven fabric is filled with resin powder and heat treated to impart enhanced strength and water repellency.

The resin powder may be filled into the carbon fiber
10 woven fabric of the invention in a state of dispersion such as in a dispersion-type solution. The filling method may be coating, an immersion or the like.

The resin powder-filled carbon fiber woven fabric is heat treated at the melting temperature of the resin.
15 Heating at the melting temperature is the optimum temperature for uniform penetration into the woven fabric as the powder resin undergoes a reduction in viscosity with increasing temperature. If the heating temperature is increased too high, deterioration of the resin or
20 foaming of the resin may occur, resulting in a lack of strength-enhancing effect. Foaming may also reduce the gas permeability. Consequently, it is preferred to conduct the heating at a temperature of no more than about 100°C higher than the softening temperature of the
25 resin. If the heating temperature is too low, on the other hand, the viscosity of the resin is not sufficiently lowered, and the filling is inadequate. The heating is therefore preferably conducted at a temperature of at least 50°C higher than the softening
30 temperature.

The carbon fiber woven fabric of the invention obtained in the manner described above is useful as a porous carbon sheet, catalyst sheet or the like for fuel cells.

35 Fig. 2 is an illustration of a carbon fiber woven fabric of the invention used as a gas diffusing porous carbon sheet for a solid polymer fuel cell.

In Fig. 2, a porous anode gas diffusing sheet 2 and grooved separator sheet 1 are situated at the anode side of an ion-exchange membrane 4 made of a solid polymer electrolyte, via an anode catalyst layer 3 made of Pt-carrying carbon black or the like, while a porous cathode gas diffusing sheet 6 and grooved separator sheet 1 are situated at the cathode side, via a cathode catalyst layer 5 made of Pt-carrying carbon black or the like. The solid polymer electrolyte 4 may be perfluorocarbonsulfonic acid, and the ion-exchange membrane has a thickness of, for example, about 50-200 μm . The grooved separator sheet 1 is fabricated from a molded expanding graphite sheet or a carbon black molded sheet, and the grooves 1a, 1b form flow channels for the fuel gas such as H_2 , the oxidizing gas, and the reaction water.

For a fuel cell construction, the gas diffusing sheets 2,6 are between the catalyst layers 3,5 and the grooved separator sheet 1, and the use of porous carbon sheets has been studied in the past to meet the requirements of chemical stability in oxidation-reduction atmospheres, high electric conductivity, high gas permeability and high compressive strength. The carbon fiber woven fabric of the invention satisfies all of the aforementioned required properties and is therefore particularly useful as a porous carbon sheet for such gas diffusing sheets.

A solid polymer fuel cell is operated at around 100°C , but the carbon fiber woven fabric of the invention can also be utilized in phosphoric acid-type fuel cells that are operated at high temperatures.

Examples

The present invention will now be explained in further detail by way of examples. The various properties in the examples were measured by the methods described below.

The thickness was measured with a micrometer.

The bulk density was calculated from the thickness and dimensions as measured with a slide caliper, and the weight as measured by a scale.

5 The compressive strength was determined by laminating 50 samples, compressing them at a loading rate of 0.5 mm/min, measuring the maximum load resistance, dividing this by the area, and calculating the maximum load resistance per unit area.

10 The layer transverse electrical resistance (layer transverse resistance) was determined by cutting out a 2 cm square (4 cm²) sample, sandwiching it between copper plates and running a direct current of 1 A/cm² through the copper plates while clamping with a load of 4 kgf/cm², and measuring the drop in voltage (V) due to
15 contact between the sample and the copper plates. The results were used to determine the flatwise electrical resistance according to the following equation.

Layer transverse electrical resistance = voltage drop (V)/current density (A/cm²)

20 The volume resistivity in the layer direction was measured by the standard four-point method.

The gas permeability was determined by passing air through a 50 cm² section of the sample at room temperature at 3000 mL/min, and calculating the
25 permeation per cm² per hour during that time from the pressure loss (PmmAq), according to the equation given below.

Gas permeability = 3000 mL/min × 60 min/50 cm²/PmmAq

30 The orientation was measured by q/(p+q) as defined above, based on SEM observation. Since all of the fabrics in Tables 1 to 3 are plain weave fabrics with uniform compositions, the orientations represent both the orientations of the specific orienting components and the average orientations.

35 The water repellency was determined by cutting out a short strip with a width of 15 mm and a length of 25 mm from the sample, immersing 5 mm of the tip in distilled

00013641-084004

the water in cm units from the position in the water to the position to which the water was drawn up, and evaluating the water repellency based on the condition of water absorption.

5 Examples 1-4

10 The following examples employed a plain woven cotton cloth (hereunder referred to as "cotton cloth A") with spun cotton yarn of 200 μ m thickness using a conventional fiber size of 20 μ m with a fabric thickness of 0.3 mm, and aluminum phosphate.

15 The meter yarn count of the woven fabric yarn was No. 20 (20 km/kg), and the plain weave thread count was 60 x 60/(2.54 cm)² (specific weight: 153 g/m²).

20 Cotton cloth A was cut into ten 400 mm squares and these were used as green samples. After immersing each of these in a 15% aqueous aluminum phosphate solution, they were drawn up and drained, and used as 5 mass% aluminum phosphate soaked sheets. The soaked sheets were sandwiched between graphite wafers, embedded in packing coke and placed in an electric furnace with a non-oxidizing atmosphere for one week for carbonization firing of the ten sheets at 900°C.

25 Five of the sheets were sandwiched between graphite wafers and fired at 1800°C in an Atchison graphitizing furnace.

30 The sheets obtained after firing at 900°C were carbon fiber woven fabrics with satisfactory appearance, a thickness of 0.28 mm and 352 mm square dimensions, remaining in the shape of the plain woven green samples.

35 Similarly the sheets after firing at 1800°C were carbon fiber woven fabrics with satisfactory appearance, a thickness of 0.26 mm and 348 mm square dimensions, remaining in the shape of the plain woven green samples.

Five each of the 900°C fired samples and the 1800°C fired samples were used as property measurement samples and washed and dried, and the properties were measured

5

10

Green samples were also prepared without aluminum phosphate treatment and ten samples each fired at 900°C and 1800°C in the same manner as above. The properties of the obtained carbon fiber woven fabrics were measured in the same manner as Examples 1-2. The results for Examples 3-4 are also shown in Table 1.

Green samples were also prepared without aluminum phosphate treatment and ten samples each fired at 900°C and 1800°C in the same manner as above. The properties of the obtained carbon fiber woven fabrics were measured in the same manner as Examples 1-2. The results for Examples 3-4 are also shown in Table 1.

Table 1

	Pretreatment	Firing temperature °C	Fabric thickness mm	Bulk density g/cm ³	Compressive strength kgf/cm ²	Layer transverse resistance mΩ · cm ²	Gas permeability cc/cm ² /hr/mmAq	Resistivity Ω · cm	Orientation
Example 1	aluminum phosphate	900	0.28-0.29	0.42-0.43	73-78	21-23	7700-8100	0.13-0.15	≥1/3
Example 2	aluminum phosphate	1800	0.26-0.26	0.44-0.44	92-97	18-20	4100-4500	0.12-0.14	≥1/3
Example 3	none	900	0.23-0.23	0.28-0.29	30-35	43-48	8500-9100	0.12-0.14	≥1/3
Example 4	none	1800	0.21-0.21	0.30-0.30	41-45	40-44	4500-5100	0.11-0.13	≥1/3

Examples 5-8

Four of the carbon fiber woven fabrics from each of Examples 1-4 were immersed in a Teflon dispersion solution (FEP Dispersion Solution, Daikin Industries Co., Ltd.) and drawn out from the solution, to fill each carbon fiber woven fabric with the FEP powder to 10 mass%.

Each of these was sandwiched between graphite wafers and heat treated at 300°C for 30 minutes. After the heat treatment, the compressive strength, layer transverse resistance and gas permeability were measured in the same manner as Examples 1-4. The results for Examples 5-8 are shown in Table 2.

The results demonstrate that addition of Teflon enhances the compressive strength without significantly lowering performance in terms of layer transverse resistance or gas permeability.

Examples 11-14

One each of the remaining carbon fiber woven fabrics of Examples 1-4 was also subjected to property measurements. The results for Examples 11-14 are also shown in Table 2. This table demonstrates that addition of Teflon prevents water absorption and enhances water repellency.

Table 2

	Pretreatment	Firing temperature °C	Post-treatment	Compressive strength kgf/cm ²	Layer transverse resistance mΩ·cm ²	Water absorption height cm	Gas permeability cc/cm ² /hr/mmAq	Resistivity Ω·cm	Orientation
Example 5	aluminum phosphate	900	Teflon	99-108	21-37	0	6900-7400	0.13-0.15	≥1/3
Example 6	aluminum phosphate	1800	Teflon	107-118	19-22	0	3600-4000	0.13-0.16	≥1/3
Example 7	none	900	Teflon	71-77	45-50	0	7600-8100	0.13-0.14	≥1/3
Example 8	none	1800	Teflon	78-85	41-46	0	4100-4700	0.11-0.13	≥1/3
Example 11	aluminum phosphate	900	none	77	21	4	7100	0.13-0.15	≥1/3
Example 12	aluminum phosphate	1800	none	95	20	3	4400	0.11-0.13	≥1/3
Example 13	none	900	none	33	48	3	8900	0.12-0.14	≥1/3
Example 14	none	1800	none	43	42	2	4700	0.11-0.13	≥1/3

Examples 21-32

5 A cotton cloth of the same weight per square meter
obtained by plain weaving yarns made by twisting two
yarns of the same size as the yarn of cotton cloth A used
in Examples 1 and 2 (same size of the fibers composing
the yarns) (cotton cloth B, weight: 325 g/m²) and a
cotton cloth of the same weight per square meter obtained
by plain weaving of non-twisted yarn narrower than cotton
cloth A and of a size such that the fiber size after
10 firing in a non-oxidizing atmosphere was 8 μm (cotton
cloth C), were fired at 900°C and 1800°C in the same
manner as Examples 1 and 2.

15 After firing, carbon fiber woven fabrics from cotton
cloths B and C were immersed in a dispersion solution in
the same manner as Examples 5 to 8 and then heated and
Teflon treated under the same conditions.

20 Separately, cotton cloth B was fired in a non-
oxidizing atmosphere at 900°C and 1800°C and directly
used as a carbon fiber woven fabric without Teflon
treatment, and the properties of both were examined by
the same methods as Examples 1-8. The water absorption
height was only determined for the Teflon-treated sample.

25 The results for Examples 21-32 are shown in Table 3.
The obtained carbon fiber woven fabrics had satisfactory
appearances with thicknesses of 0.2-0.3 mm.

Table 3

	Fabric type	Pre-treatment	Firing temperature °C	Post-treatment	Compressive strength kgf/cm ²	Layer transverse resistance mΩ·cm ²	Gas permeability cc/cm ² /hr/mmHg	Water absorption height	Resistivity Ω·cm	Orientation
Example 21	cotton fabric twisted yarn	none	900	Teflon	105-116	22-35	5000-5400	0	0.12-0.14	≥1/3
Example 22	cotton fabric twisted yarn	none	1800	Teflon	113-125	18-23	3800-4100	0	0.10-0.11	≥1/3
Example 23	cotton fabric narrow yarn	none	900	Teflon	103-115	24-36	4500-4900	0	0.13-0.15	≥1/3
Example 24	cotton fabric narrow yarn	none	1800	Teflon	110-121	17-22	3300-3600	0	0.12-0.14	≥1/3
Example 25	cotton fabric twisted yarn	none	900	none	71-73	44-48	6000-6300	-	0.12-0.14	≥1/3
Example 26	cotton fabric twisted yarn	none	1800	none	87-90	39-43	4400-4800	-	0.10-0.11	≥1/3
Example 27	cotton fabric narrow yarn	none	900	none	70-73	40-44	5800-6100	-	0.12-0.13	≥1/3
Example 28	cotton fabric narrow yarn	none	1800	none	80-85	38-41	4200-4400	-	0.11-0.12	≥1/3
Example 29	cotton fabric twisted yarn	none	2200	none	82-88	18-25	2800-3200	-	0.11-0.11	≥1/3
Example 30	cotton fabric twisted yarn	none	2600	none	70-76	15-22	2900-3500	-	0.09-0.09	≥1/3
Example 31	cotton fabric twisted yarn	none	2200	Teflon	110-125	23-29	2700-3300	0	0.12-0.13	≥1/3
Example 32	cotton fabric twisted yarn	none	2600	Teflon	105-118	17-26	2800-3400	0	0.10-0.11	≥1/3

These results demonstrate that these obtained carbon fiber woven fabrics were also carbon fiber woven fabrics with excellent properties, that the same properties are obtained even when cotton cloths woven with twisted or narrow yarn are fired at a prescribed temperature in a non-oxidizing atmosphere, and especially that Teflon treatment imparts water repellency and enhances the strength properties.

10 Industrial Applicability

15 The carbon fiber woven fabric according to the invention can be obtained by a low cost process suitable for mass production, using a cheap and readily available commercial cellulose-based fiber woven fabric as the starting material, and since the carbon fiber woven fabric is obtained in a form retaining the shape of the starting fabric, it is easy to handle and is useful as a gas diffusing porous carbon sheet, catalyst sheet or the like for alkali-type, phosphoric acid-type or solid polymer-type fuel cells which require the use of carbon sheet materials.

CLAIMS

1. A carbon fiber woven fabric characterized by being obtained by firing a cellulose-based woven fabric, and by having a thickness in the range of 0.05-0.4 mm, a
5 volume resistivity of not less than $0.2 \Omega \cdot \text{cm}$ in the layer direction, and a gas permeability of not less than $1500 \text{ cc/cm}^2/\text{hr/mmHg}$.

2. The carbon fiber woven fabric as claimed in claim 1, wherein the compressive strength is not less
10 than 70 kgf/cm^2 .

3. The carbon fiber woven fabric as claimed in claim 1, wherein the electrical resistance in the direction of thickness of the woven fabric is no greater than $50 \text{ m}\Omega \cdot \text{cm}^2$ as measured between two copper plates with
15 a load of 4 kgf/cm^2 .

4. The carbon fiber woven fabric as claimed in claim 1, wherein the orientation of the carbon fiber woven fabric as defined in the present specification includes an orientation component of $4/9$ or greater.

20 5. The carbon fiber woven fabric as claimed in claim 1, wherein the orientation of the carbon fiber woven fabric as defined in the present specification is an average of $1/3$ or greater.

25 6. The carbon fiber woven fabric as claimed in claim 1 which is a plain weave.

7. The carbon fiber woven fabric as claimed in claim 1 which has a water repellent property.

30 8. The gas diffusion porous carbon sheet for a solid polymer fuel cell which comprises a carbon fiber woven fabric as claimed in claim 1.

9. (Amended) A process for manufacture of a carbon fiber woven fabric as claimed in claim 1, characterized by firing a cellulose-based woven fabric in a non-oxidizing atmosphere.

35 10. The process for manufacture of a carbon fiber woven fabric as claimed in claim 9, wherein said

09913611-081501
FOOTED T 961501

- 25/1 -

cellulose-based woven fabric is soaked with a phosphoric

0913611 001601
TOP SECRET

11. The process for the manufacture of a carbon fiber woven fabric as claimed in claim 9, wherein the firing temperature is in the range of 900-3000°C.

5

Fig.1B

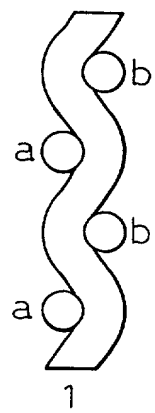


Fig.1A

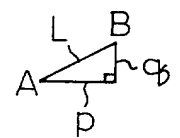
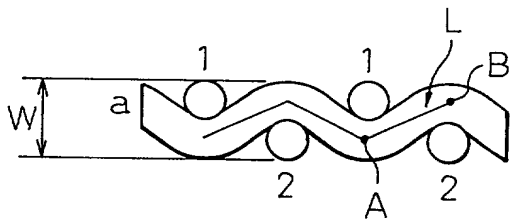
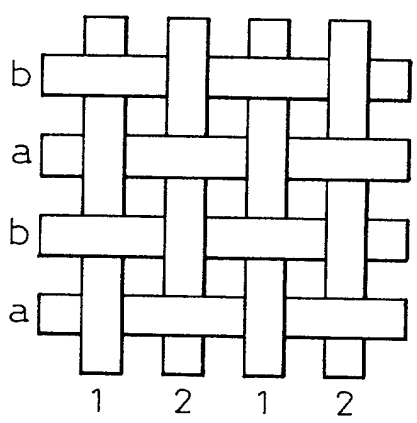
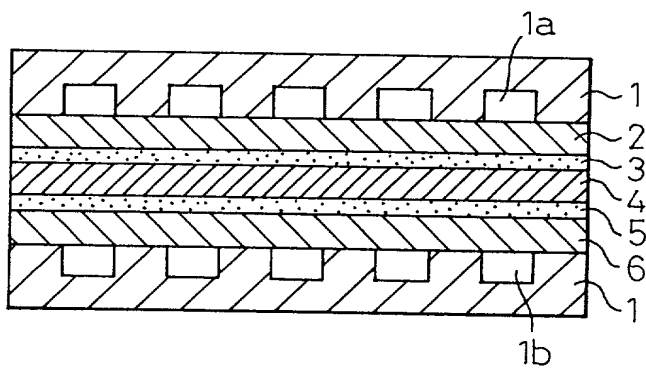


Fig.1C

Fig.2



Declaration and Power of Attorney for Patent Application

特許出願宣言書及び委任状

Japanese Language Declaration

日本語宣言書

下記の氏名の発明者として、私は以下の通り宣言します。

As a below named inventor, I hereby declare that:

私の住所、私書箱、国籍は下記の私の氏名の後に記載された通りです。

My residence, post office address and citizenship are as stated next to my name.

下記の名称の発明に関して請求範囲に記載され、特許出願している発明内容について、私が最初かつ唯一の発明者（下記の氏名が一つの場合）もしくは最初かつ共同発明者であると（下記の名称が複数の場合）信じています。

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled

CARBON FIBER WOVEN FABRIC
AND METHOD FOR
PRODUCTION THEREOF

上記発明の明細書（下記の欄でx印がついていない場合は、本書に添付）は、

the specification of which is attached hereto unless the following box is checked:

- ☐ 月 日に提出され、米国出願番号または特許協定条約国際出願番号を _____ とし、
（該当する場合） _____ に訂正されました。

- ☐ was filed on February 16, 2000
as United States Application Number or PCT
International Application Number PCT/JP00/00875
and was amended on December 21, 2000
[B2] (if applicable)

(Under PCT Article 34)

私は、特許請求範囲を含む上記訂正後の明細書を検討し、内容を理解していることをここに表明します。

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above.

私は、連邦規則法典第37編第1条56項に定義されるとおり、特許資格の有無について重要な情報を開示する義務があることを認めます。

I acknowledge the duty to disclose information which is material to patentability as defined in Title 37, Code of Federal Regulations, § 1.56.

Japanese Language Declaration

日本語宣言書

私は、米国法典第35編119条(a)-(d)項又は365条(b)項に基づき下記の、米国外の国の少なくとも一カ国を指定している特許協力条約365(a)項に基づき国際出願、又は外国での特許出願もしくは発明者証の出願についての外国優先権をここに主張するとともに、優先権を主張している。本出願の前に出願された特許または発明者証の外国出願を以下に、枠内をマークすることで、示しています。

Prior foreign application(s)

外国での先行出願
11-40518

[C1] (Pat. Appln.)

(Number)

(番号)

11-161756

[C5] (Pat. Appln.)

(Number)

(番号)

[C9]

(Number)

(番号)

[C2] Japan

(Country)

(国名)

[C6] Japan

(Country)

(国名)

[C10]

(Country)

(国名)

私は、第35編米国法典119条(e)項に基づいて下記の米
国特許出願規定に記載された権利をここに主張いたします。

[D1]

(Application No.)

(出願番号)

[D2]

(Filing Date)

(出願日)

私は、下記の米国法典第35編120条に基づいて下記の米
国特許出願に記載された権利、又は米国を指定している特許
協力条約365条(c)に基づき権利をここに主張します。ま
た、本出願の各請求範囲の内容が米国法典第35編112条
第1項又は特許協力条約で規定された方法で先行する米国特
許出願に開示されていない限り、その先行米国出願書提出日
以降で本出願書の日本国内または特許協力条約国際提出日ま
での期間中に入手された、連邦規則法典第37編1条56項
で定義された特許資格の有無に関する重要な情報について開
示義務があることを認識しています。

[E1]

(Application No.)

(出願番号)

[E2]

(Filing Date)

(出願日)

[E4]

(Application No.)

(出願番号)

[E5]

(Filing Date)

(出願日)

私は、私自身の知識に基づいて本宣言書中で私が行なう表
明が真実であり、かつ私の入手した情報と私の信じることに
基づき表明が全て真実であると信じていること、さらに故
意になされた虚偽の表明及びそれと同等の行為は米国法典第
18編第1001条に基づき、罰金または拘禁、もしくはその
両方により処罰されること、そしてそのような故意による
虚偽の声明を行えば、出願した、又は既に許可された特許
の有効性が失われることを認識し、よってここに上記のごと
く宣誓を致します。

I hereby claim foreign priority under Title 35, United States Code, § 119(a)-(d) or § 365 (b) of any foreign application(s) for patent or inventor's certificate, or § 365(a) of any PCT International application which designated at least one country other than the United States, listed below and have also identified below, by checking the box, any foreign application for patent or inventor's certificate, or PCT International application having a filing date before that of the application on which priority is claimed.

Priority Not Claimed

優先権主張なし

[C3] 18/February/1999

(Day/Month/Year Filed)

(出願年月日)

[C7] 9/June/1999

(Day/Month/Year Filed)

(出願年月日)

[C11]

(Day/Month/Year Filed)

(出願年月日)

I hereby claim the benefit under Title 35, United States Code, § 119(e) of any United States provisional application(s) listed below. 60/128,054 7/April/1999

[D3] 60/141,244

(Application No.)

(出願番号)

(both of Pat. Applns.)

[D4] 30/June/1999

(Filing Date)

(出願日)

I hereby claim the benefit under Title 35, United States Code, § 120 of any United States application(s), or § 365(c) of any PCT International application designating the United States, listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States or PCT International application in the manner provided by the first paragraph of Title 35, United States Code, § 112, I acknowledge the duty to disclose information which is material to patentability as defined in Title 37, Code of Federal Regulations, § 1.56 which became available between the filing date of the prior application and the national or PCT International filing date of this application.

[E3]

(Status)(patented, pending, abandoned)

(現況：特許許可済、保属中、放棄済)

[E6]

(Status)(patented, pending, abandoned)

(現況：特許許可済、保属中、放棄済)

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Japanese Language Declaration

日本語宣言書

委任状： 私は下記の発明者として、本出願に関する一切の手続きを米特許商標局に対して遂行する弁理士または代理人として、下記の者を指名いたします。（弁理士、または代理人の氏名及び登録番号を明記のこと）

POWER OF ATTORNEY: As a named inventor, I hereby appoint the following attorney(s) and/or agent(s) to prosecute this application and transact all business in the Patent and Trademark Office connected therewith: (list name and registration number)

John H. Mion, Reg. No. 18,879; Thomas J. Macpeak, Reg. No. 19,292; Robert J. Seas, Jr., Reg. No. 21,092; Darryl Mexic, Reg. No. 23,063; Robert V. Sloan, Reg. No. 22,775; Peter D. Olexy, Reg. No. 24,513; J. Frank Osha, Reg. No. 24,625; Waddell A. Biggart, Reg. No. 24,861; Louis Gubinsky, Reg. No. 24,835; Neil B. Siegel, Reg. No. 25,200; David J. Cushing, Reg. No. 28,703; John R. Inge, Reg. No. 26,916; Joseph J. Ruch, Jr., Reg. No. 26,577; Sheldon I. Landsman, Reg. No. 25,430; Richard C. Turner, Reg. No. 29,710; Howard L. Bernstein, Reg. No. 25,665; Alan J. Kasper, Reg. No. 25,426; Kenneth J. Burchfiel, Reg. No. 31,333; Gordon Kit, Reg. No. 30,764; Susan J. Mack, Reg. No. 30,951; Frank L. Bernstein, Reg. No. 31,484; Mark Boland, Reg. No. 32,197; William H. Mandir, Reg. No. 32,156; Brian W. Hannon, Reg. No. 32,778; Abraham J. Rosner, Reg. No. 33,276; Bruce E. Kramer, Reg. No. 33,725; Paul F. Neils, Reg. No. 33,102; Brett S. Sylvester, Reg. No. 32,765; Robert M. Masters, Reg. No. 35,603; George F. Lehnigk, Reg. No. 36,359; John T. Callahan, Reg. No. 32,607 and Steven M. Gruskin, Reg. No. 36,818.

書類送付先

Send Correspondence to:
SUGHRUE, MION, ZINN, MACPEAK & SEAS, PLLC
2100 Pennsylvania Avenue, N.W. Suite 800
Washington, D.C. 20037-3213

直接電話連絡先：（名前及び電話番号）

Direct Telephone Calls to:

(202)293-7060

唯一または第一発明者名 1-00	Full name of sole or first inventor Shigeru Murakami
発明者の署名 日付	Inventor's signature Shigeru Murakami Date August 6, 2001
住所	Residence Omachi-shi, Nagano, Japan JPY
国籍	Citizenship Japanese
私書箱	Post Office Address c/o SHOWA DENKO K.K. OMACHI WORKS, 6850, Oaza Omachi, Omachi-shi, Nagano 399-6461, Japan
第二共同発明者 2-00	Full name of second joint inventor, if any Yoichi Namba
第二共同発明者 日付	Second inventor's signature Yoichi Namba Date August 6, 2001
住所	Residence Omachi-shi, Nagano, Japan JPY
国籍	Citizenship Japanese
私書箱	Post Office Address c/o SHOWA DENKO K.K. OMACHI WORKS, 6850, Oaza Omachi, Omachi-shi, Nagano 399-6461, Japan

（第三以降の共同発明者についても同様に記載し、署名をすること）

(Supply similar information and signature for third and subsequent joint inventors.)

Japanese Language Declaration

日本語宣言書

第三の共同発明者の氏名 (該当する場合)		3-00	Full name of third joint inventor, if any <u>Tsutomu Masuko</u>
同第三発明者の署名	日付		Third inventor's signature <u>Tsutomu Masuko</u> Date August 6, 2001
住所			Residence <u>Omachi-shi, Nagano, Japan</u>
国籍			Citizenship <u>Japanese</u> JPX
郵便の宛先			Post office address <u>c/o SHOWA DENKO K.K., OMACHI WORKS,</u>
			<u>6850, Oaza Omachi, Omachi-shi,</u>
			<u>Nagano 399-6461, Japan</u>
第四の共同発明者の氏名 (該当する場合)		4-00	Full name of fourth joint inventor, if any <u>Yuichi Kamijo</u>
同第四発明者の署名	日付		Fourth inventor's signature <u>Yuichi Kamijo</u> Date August 6, 2001
住所			Residence <u>Omachi-shi, Nagano, Japan</u>
国籍			Citizenship <u>Japanese</u> JPX
郵便の宛先			Post office address <u>c/o SHOWA DENKO K.K. OMACHI WORKS,</u>
			<u>6850, Oaza Omachi, Omachi-shi,</u>
			<u>Nagano 399-6461, Japan</u>
第五の共同発明者の氏名 (該当する場合)			Full name of fifth joint inventor, if any
同第五発明者の署名	日付		Fifth inventor's signature Date
住所			Residence
国籍			Citizenship
郵便の宛先			Post office address
第六の共同発明者の氏名 (該当する場合)			Full name of sixth joint inventor, if any
同第六発明者の署名	日付		Sixth inventor's signature Date
住所			Residence
国籍			Citizenship
郵便の宛先			Post office address